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Variants in the melatonin receptor 1B gene (*MTNR1B*) influence fasting glucose levels

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Abstract

To identify novel genetic loci associated with fasting glucose concentrations, we examined the leading association signals in 10 genome-wide association scans involving a total of 36,610 individuals of European descent. Variants in the gene encoding the melatonin receptor 1B (*MTNR1B*) were consistently associated with fasting glucose across all ten studies. The strongest signal was observed at rs10830963, where each G-allele (frequency 0.30 in HapMap CEU) was associated with an increase of 0.07 (95%CI 0.06–0.08) mmol/L in fasting glucose levels ($P=3.2\times 10^{-50}$) and reduced beta-cell function as measured by homeostasis model assessment (HOMA-B, $P=1.1\times 10^{-15}$). The same allele was associated with an increased risk of type 2 diabetes (odds ratio = 1.09 (1.05–1.12), per G allele $P=3.3\times 10^{-7}$) in a meta-analysis of thirteen case-control studies totalling 18,236 cases and 64,453 controls. Our analyses also confirm previous associations of fasting glucose with variants at the *G6PC2* (rs560887, $P=1.1\times 10^{-57}$) and *GCK* (rs4607517, $P=1.0\times 10^{-25}$) loci.

Blood and plasma fasting glucose (FG) levels are usually tightly regulated within a narrow physiologic range by a feedback mechanism that targets a particular FG set point for each individual^{1,2}. Disruption of normal glucose homeostasis and substantial elevations of FG are hallmarks of type 2 diabetes (T2D) and typically result from sustained reduction in pancreatic beta-cell function and insulin secretion.

However, even within healthy, non-diabetic populations there is substantial variation in FG levels. Approximately one-third of this variation is genetic³, but little of this heritability has been explained. There is growing evidence to suggest that common variants contributing to variation in FG are largely distinct from those associated with major disruptions of beta-cell function that predispose to T2D. Common sequence variants in the glucokinase (*GCK*) promoter^{4–6}, and around genes encoding the islet specific glucose-6-phosphatase (*G6PC2*)^{5,6} and the glucokinase regulatory protein (*GCKR*)^{7–9} have each been associated with individual variation in FG levels, but have, at best, weak effects on T2D risk^{8,10}. Furthermore, though there are now over 15 genetic loci strongly associated with the risk of T2D^{7,10–14}, none shows compelling evidence for association with FG in the two genome-wide association scans (GWAS) so far reported^{5,6}.

MAGIC (the Meta-Analyses of Glucose and Insulin-related traits Consortium) represents a collaborative effort to combine data from multiple GWAS to identify additional loci that impact on glycaemic and metabolic traits. Our genetic studies of FG levels originally coalesced into four distinct consortia: (i) European Network for Genetic and Genomic Epidemiology (ENGAGE), combining data from deCODE, Northern Finland Birth Cohort 1966 (NFBC1966), Netherlands Twins Register/Netherlands Study of Depression and Anxiety (NTR/NESDA), and the Rotterdam Study; (ii) Genetics of Energy Metabolism (GEM), a meta-analysis of the Lausanne (CoLaus) and TwinsUK scans; (iii) DFS, involving the Diabetes

Genetics Initiative (DGI), Finland-United States Investigation of NIDDM Genetics (FUSION) and SardinIA scans; and (iv) the Framingham Heart Study (FHS). Details of the ten component studies (n=1,233–6,479) are provided in Supplementary Table 1.

As a prelude to more extensive data-sharing, the four consortia initially exchanged the identities of between 10 and 20 SNPs prominently associated with FG in their individual, interim, meta-analyses (n=6,479–12,389; Supplementary Table 2). Comparison of these signals revealed three loci with consistent effects on FG detected in multiple studies. Two of these represented the previously reported signals in *G6PC2* and *GCK*. In addition, all four groups independently generated evidence for an association between FG and SNPs around the *MTNR1B* (melatonin receptor 1B) locus (ENGAGE [rs1387153] $P=2.2 \times 10^{-17}$; GEM [rs10830963] $P=7.4 \times 10^{-11}$; DFS [rs10830963] $P=2.5 \times 10^{-7}$; FHS [rs11020107] $P=5.8 \times 10^{-4}$, for the most strongly associated SNP exchanged from each analysis). The association signals at all three loci were confirmed on formal meta-analysis including results from all 10 studies, after exclusion of individuals with known diabetes (rs560887 [*G6PC2*], $P=1.1 \times 10^{-57}$; rs4607517 [*GCK*], $P=1.0 \times 10^{-25}$; rs10830963 [*MTNR1B*], $P=3.2 \times 10^{-50}$) (Table 1, Supplementary table 3). Subsequent efforts to harmonize additional aspects of data analysis strategies (including the additional exclusion, where necessary, of individuals with FG measures >7mmol/l) had only a marginal impact on estimates of significance and effect size (Supplementary Table 4).

We attempted to refine the location of the *MTNR1B* association signal by extending the meta-analysis to all SNPs (genotyped and imputed from the HapMap) within the 1Mb region flanking the gene (n=35,812; 981 SNPs). In all, 30 genotyped and imputed SNPs showed compelling evidence for association with FG ($P<10^{-8}$). The strongest signal was detected at rs10830963: the minor (G) allele (frequency 0.30 in HapMap CEU¹⁵) at this SNP was associated with a per-allele increase of 0.07 (95%CI 0.06–0.08) mmol/L in FG ($P=3.2 \times 10^{-50}$). Consistent evidence for association at rs10830963 was observed in all 10 component GWAS, irrespective of whether this SNP was genotyped or imputed, and the genotyping platform (Table 1, Supplementary Table 1). Repeat meta-analysis within the region after conditioning on rs10830963 revealed no additional independent signals of association (Supplementary Note).

The strength of the association between rs10830963 and FG was unchanged after adjustment for body mass index (Supplementary Table 4). Analyses of fasting insulin levels as well as indices of beta-cell function (HOMA-B) and insulin sensitivity (HOMA-IR) estimated by the homeostasis model assessment¹⁶ were possible in ~24,000 participants from the 10 studies. These established that the glucose-raising allele at rs10830963 was associated with reduced beta-cell function ($P=1.1 \times 10^{-15}$), with no appreciable effect on fasting insulin or insulin sensitivity (Supplementary Table 5, Supplementary Note).

To determine the impact of variants within *MTNR1B* on T2D risk, we performed a large-scale meta-analysis of thirteen T2D case-control samples (18,236 T2D cases, 64,453 controls; corresponding to an effective sample size of 21,179 unrelated cases and 21,179 unrelated controls). We combined data from the deCODE¹³, Rotterdam¹⁷, KORA¹⁸, FUSION Stage 2¹¹ and METSIM¹⁰ studies and from several case-control samples from the UK¹⁰ with publicly-available data from the DIAGRAM consortium (which itself aggregates GWA data from the WTCCC, DGI and FUSION scans)¹⁰ (Supplementary Note). We found strong evidence that the minor G-allele of rs10830963 was associated with increased risk of T2D (odds ratio=1.09 [1.05–1.12], $P=3.3 \times 10^{-7}$) (Supplementary Table 6 and Figure 2). The possibility that the FG association might reflect the inclusion within the cross-sectional study samples of subjects with undiagnosed T2D can be discounted given that exclusion of those with either known diabetes, or a FG exceeding 7mmol/l had little impact on the strength of the association signal (Table 1· Supplementary Table 4). Although the association with T2D does not, despite large-scale replication efforts, reach the 5×10^{-8} threshold consistent with

“genome-wide significance”¹⁵, it seems highly probable, given the strong impact of this variant on beta-cell function (Supplementary Table 5), that this is a genuine effect.

The analyses we performed interrogate only a minority of common sequence variants in a given region – it is likely that the causal variant for this locus is yet to be identified. The SNP with the strongest statistical evidence so far, rs10830963, maps within the single 11.5 kb intron of *MTNR1B* but does not appear to disrupt consensus transcription factor binding or cryptic alternative splice sites. The association signal is bounded by recombination hotspots defining a ~60kb interval within which all our strongly associated SNPs lie and the causal variant is likely to reside. This interval contains the entire coding region of *MTNR1B*. The only other nearby genes (the coding regions of which lie well outside this 60kb region) are *SLC36A4* and *FAT3*, neither of which are compelling candidates. *SLC36A4* encodes a proton/amino acid transmembrane transporter moderately similar to *Rattus norvegicus* lysosomal amino acid transporter 1, while *FAT3* encodes a cadherin family member which is the human homolog of the *Drosophila melanogaster* FAT tumour suppressor gene. Ultimately, detailed fine-mapping and functional analyses will be required to define the causal allele(s) and to confirm that this effect is mediated through altered function or expression of *MTNR1B*.

As well as exploring this novel signal, the size of the MAGIC data-set allowed us to examine the *G6PC2* and *GCK* regions in greater detail than has previously been possible. In the *G6PC2* region, rs560887, within intron 3 of the gene, remained the strongest signal whether or not imputed data were included ($P=1.1 \times 10^{-57}$ across all 10 studies: Supplementary Figure 1). This is the same SNP reported in one recent paper,⁵ and in substantial LD ($r^2=0.72$ in HapMap CEU) with the lead SNP (rs563694) identified in a second⁶. In the *GCK* region, rs4607517, which lies 6.6 kb upstream of the gene, was the most strongly-associated SNP ($P=1.0 \times 10^{-25}$) (Supplementary Figure 1· Table 1). This SNP is also in strong LD ($r^2 = 1$ in HapMap CEU) with the *GCK* promoter SNP (rs1799884) that was featured in previous reports⁴. Repeat meta-analysis after conditioning on the respective lead SNPs revealed no additional independent association signals at either locus (Supplementary Note).

As with *MTNR1B*, the magnitudes of the FG associations for both these signals were unchanged after adjustment for BMI (Supplementary Table 4). Glucose-raising alleles at *GCK* and *G6PC2* were associated with reduced beta-cell function (rs4607517A, $P=9.8 \times 10^{-6}$; rs560887C, $P=1.2 \times 10^{-26}$) (Supplementary Table 5, Supplementary Note). However, in line with previous reports^{4,9}, neither signal was strongly associated with T2D in the large-scale meta-analysis: in fact, the glucose-raising allele at *G6PC2* was weakly associated with reduced T2D risk (rs4607517A, per-allele OR 1.05 [1.00–1.10], $P=0.031$; rs560887C, 0.93 [0.89–0.97], $P=0.0017$) (Supplementary Table 6).

We found no influence of the non-coding lead SNPs rs10830963, rs560887 or rs4607517 on gene expression of *MTNR1B*, *SLC36A4*, *FAT3*, *G6PC2* or *GCK* in genome-wide expression QTL datasets from lymphocyte derived cell lines^{19,20}, cerebral cortex²¹, or liver²², and no evidence for epistatic effects among the three lead SNPs was observed (P -2 way interactions > 0.19 in each of the 7 studies including only unrelated individuals; interactions were not examined in the other 3 studies).

MTNR1B encodes one of two known human melatonin receptors²³. Although this is the first study to implicate genetic variation in *MTNR1B* in the regulation of FG levels and predisposition to T2D, this relationship is biologically credible. As well as being highly-expressed in the brain, retina and elsewhere²⁴, *MTNR1B* is transcribed in human islets and rodent insulinoma cell lines²⁵ and the translated receptor is thought to mediate the inhibitory effect of melatonin on insulin secretion²⁶. Melatonin release is characterized by marked circadian variability and these inhibitory effects on insulin secretion may contribute to the

entrainment of circadian patterns of insulin release²⁷. There is substantial evidence in human and rodent studies linking disturbances of circadian rhythmicity to metabolic conditions including diabetes^{28,29} and over-expression of melatonin receptors has been observed in islets from patients with T2D as compared to non-diabetic controls³⁰. Taken together, these findings suggest that the association with raised FG and T2D may be driven by variants which augment expression and/or activity of islet melatonin receptors.

Our findings bring the number of common variant loci influencing FG levels to four, three of which were detected in the present study. Variants in *GCKR* have a smaller effect size than the others^{7,9} and the present study design (based on exchange of a limited number of prominent signals between component groups) was not well-powered to detect these. However, subsequent meta-analysis of *GCKR* variant data across all 10 study samples confirms the association with FG (rs780094, $P=8.5 \times 10^{-9}$) (Supplementary Table 4). The total variance in FG presently attributable to these four signals is 1.5%, indicating that additional loci remain to be found³. In comparison with *GCK* and *G6PC2*, variants in *MTNR1B* appear to have a more marked effect on risk of T2D, the effect size being comparable in magnitude (OR=1.09 [1.05–1.12]) to several other T2D-susceptibility genes recently identified in GWAS¹⁰. Thus, whilst the physiological regulation of FG set point and the pathological decline in beta-cell function which characterizes common forms of T2D generally appear to involve different processes, the *MTNR1B* finding suggests that this is not always the case. Not only can the study of diabetes-related quantitative traits provide an important path to the identification of additional T2D susceptibility loci, but there may also be opportunities for useful therapeutic overlap.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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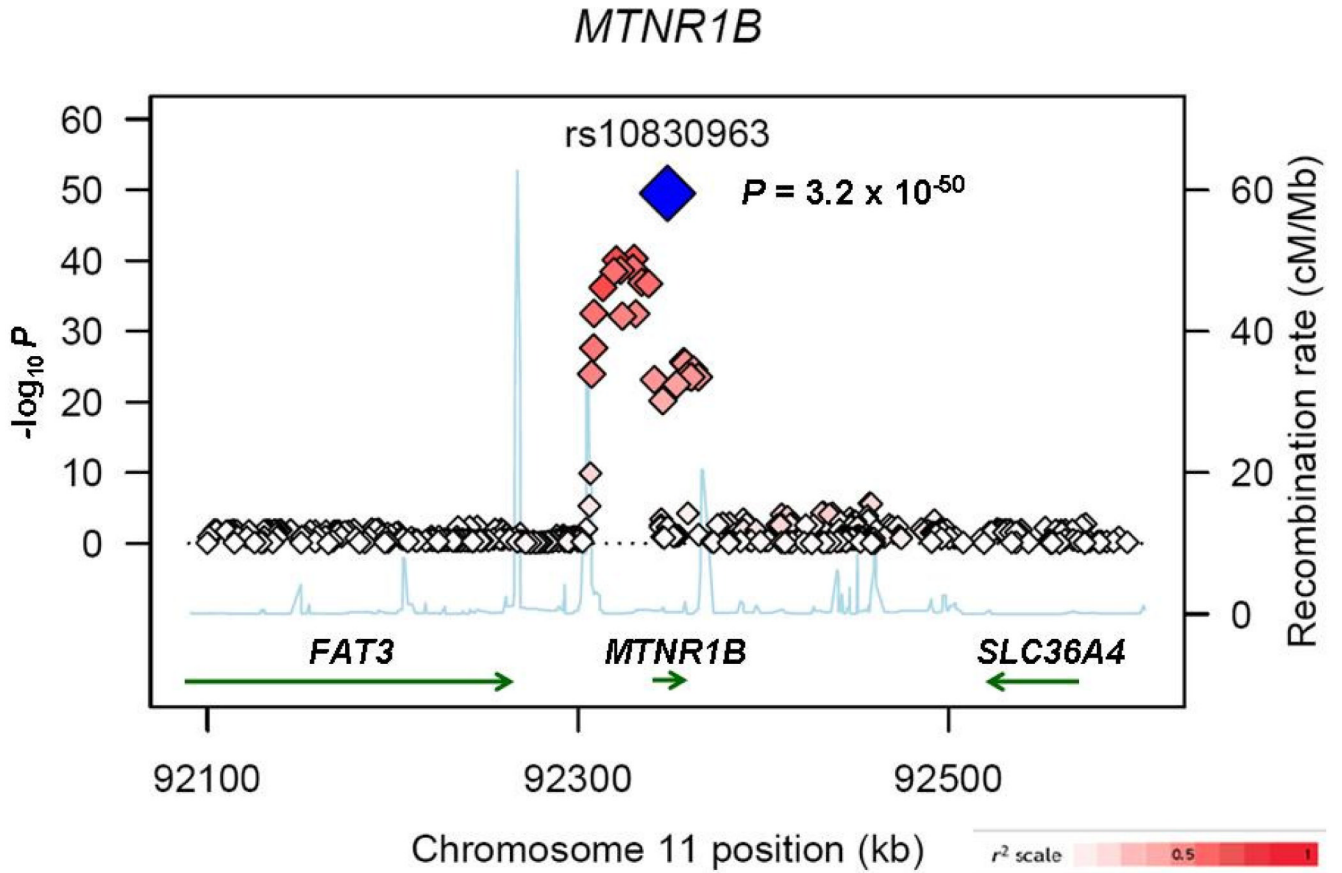


Figure 1. Regional plot of fasting glucose association results for the *MTNR1B* locus across 10 MAGIC GWAS
 Meta-analysis $-\log_{10} P$ -values are plotted as a function of genomic position (NCBI Build 35). The SNP with the strongest signal (rs10830963) is denoted by a blue diamond. Estimated recombination rates (from HapMap) are plotted to reflect the local linkage disequilibrium structure around associated SNPs and proxies (according to a white-to-red scale from $r^2=0$ to $r^2=1$; based on pair-wise r^2 values from HapMap CEU). Gene annotations were taken from the University of California-Santa Cruz genome browser.

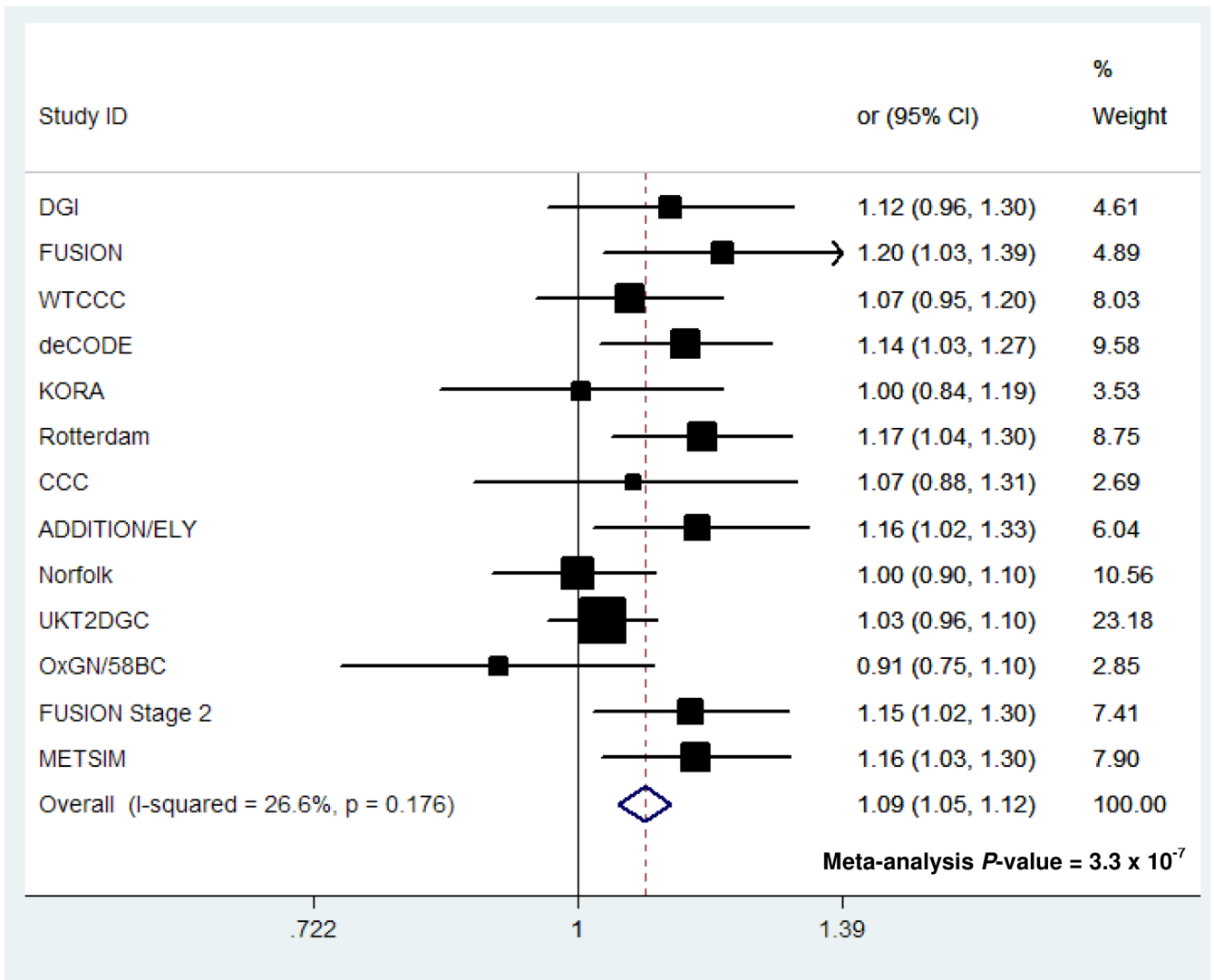


Figure 2. Association of rs10830963 with type 2 diabetes (T2D) in thirteen case-control studies

Table 1

Associates SNPs across all ten studies for three fasting glucose loci (MTNR1B, G6PC2 and GCK)

FG levels (mmol/L) are reported untransformed and unadjusted for covariates. Effect of the risk allele and SE were calculated using untransformed FG values. P-values are reported for the additive genetic model with study-specific transformation of FG values, adjusted for gender and age).

Study sample	N	G allele Frequency	Mean level of FG ^{***} per genotype (SD), mmol/L			Per_allele effect (SE), mmol/L	P-value
			CC	CG	GG		
CoL.aus	5,000	0.32	5.36 (0.71)	5.46 (0.80)	5.54 (0.81)	0.094 (0.016)	1.9 × 10 ⁻⁹
deCODE	6,240	0.27	5.29 (0.71)	5.39 (0.71)	5.44 (0.71)	0.086 (0.016)	9.2 × 10 ⁻⁸
DGI	1,455	0.31	5.29 (0.54)	5.32 (0.53)	5.39 (0.60)	0.042 (0.022)	0.054
Framingham*	6,479	0.28	5.16 (0.48)	5.21 (0.48)	5.26 (0.46)	0.050 (0.012)	2.2 × 10 ⁻¹³
FUSION	1,233	0.33	5.28 (0.49)	5.33 (0.47)	5.40 (0.44)	0.057 (0.016)	5.8 × 10 ⁻⁴
NFBC1966	4,245	0.34	5.63 (0.46)	5.70 (0.49)	5.80 (0.46)	0.079 (0.012)	1.7 × 10 ⁻¹¹
NTR/NESDA	3,166	0.27	5.22 (0.64)	5.26 (0.62)	5.38 (0.63)	0.062 (0.019)	1.2 × 10 ⁻³
Rotterdam	2,058	0.28	5.58 (0.81)	5.75 (0.91)	5.83 (1.03)	0.145 (0.029)	7.9 × 10 ⁻⁷
Sardinia	4,108	0.20	5.62 (0.89)	5.68 (0.89)	5.76 (0.89)	0.070 (0.019)	3.2 × 10 ⁻⁴
TwinsUK**	1,828	0.30	4.58 (0.65)	4.67 (0.50)	4.74 (0.57)	0.084 (0.032)	7.9 × 10 ⁻³
rs10830963 (MTNR1B)				Meta-analysis		0.072 (0.005)	3.2 × 10⁻⁵⁰
rs560887 (G6PC2)				Meta-analysis		0.064 (0.004)	1.1 × 10⁻⁵⁷
rs4607517 (GCK)				Meta-analysis		0.062 (0.007)	1.0 × 10⁻²⁵

* In Framingham study, mean FG values for the imputed SNPs are reported for proxies: rs560887 (proxy = rs73225, r²=0.96); rs4607517 (proxy = rs1799884, r²=1); rs10830963 (proxy = rs7936247, r²=0.59)

** In the TwinsUK study, mean FG values per genotype are estimated for a subset of unrelated individuals only

*** FG levels in NFBC1966 and Sardinia were measured in whole blood; in other samples measures were conducted on plasma samples. Values in the table are corrected to plasma FG using a correction factor of 1.13